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L40: Entry 1 of 2

File: USPT

Oct 29, 2002

DOCUMENT-IDENTIFIER: US 6472154 B1  
TITLE: Polymorphic repeats in human genes

## Detailed Description Paragraph Table (170):

L26953 3'UTR 4.16 2797 CAAAAA L27560 UNKNOWN 6 3438 GTT L27560 UNKNOWN 14 3658 A L27745 CDS 7 1067 AG L30117 UNKNOWN 79 652 T L31881 3'UTR 4.59 1481 CCCAG L32832 CDS 2.93 2092 GAGGAGGAGGAAGAA (SEQ ID NO:254) L32832 CDS 10 5866 CAA L32832 CDS 7.66 10261 CAG L32832 CDS 7 2985 GGC L33075 3'UTR 16 6723 A L33243 3'UTR 9 13947 TG L33477 3'UTR 10 3557 CA L34357 CDS 6 590 GGC L34408 UNKNOWN 15 706 A L35592 UNKNOWN 14 1577 AC L36140 5'UTR 24 1 T L36642 3'UTR 4 4333 CAAAA L36983 3'UTR 5.66 3368 CTC L37112 CDS 3.5 1257 CCTCAG L37198 UNKNOWN 23 7 A L38707 CDS 3.83 146 CCGGGG L38951 3'UTR 31 3653 A L38951 3'UTR 13 3736 T L38961 3'UTR 22 2283 T L39064 CDS 8.33 1504 AGC L39833 3'UTR 13 1809 A L39833 3'UTR 13 2713 T L40377 5'UTR 6.66 33 GCG L40377 5'UTR 6.33 51 GCA L40392 3'UTR 14 2209 T L40992 CDS 6 18 CAG L40992 BORDER 5.66 0 CAG L41690 CDS 6 620 GCC L41887 3'UTR 14 2046 A L41919 CDS 8 440 GGC L42025 5'UTR 5.66 1 GGC L42243 3'UTR 7 3934 GT L42243 3'UTR 16 2217 T L42243 3'UTR 15 1137 T L42243 3'UTR 12 2874 A L44505 UNKNOWN 13 309 A L46353 5'UTR 2.8 2094 CACACTCACA (SEQ ID NO:255) L46353 5'UTR 19 2401 TC L46353 5'UTR 9.5 2381 TC L46353 5'UTR 6.5 1394 TG L46353 5'UTR 13 264 A L46353 3'UTR 5 3378 TGGGG L48796 UNKNOWN 15 147 A L49169 3'UTR 6 1661 GAG L49169 3'UTR 6.5 2689 CT L49380 CDS 6 1771 GCC L76702 CDS 4.33 301 CAGCCC L76703 3'UTR 12 2219 A L77864 CDS 5.66 571 GAG L78833 3'UTR 18 6465 A M10901 3'UTR 18 3217 A M11220 3'UTR 5.5 693 TATT M11353 3'UTR 16 817 A M11722 5'UTR 14 216 G M12783 UNKNOWN 3.85 296 CGCAGCT M12783 UNKNOWN 7.5 3612 AC M12783 UNKNOWN 16 238 A M13232 3'UTR 6.5 1889 CA M13452 3'UTR 8.5 2030 GA M13452 3'UTR 15 1745 A M13903 CDS 5 528 GAGCAGCAGGAGGGGAGCTGGAGCTCCCA (SEQ ID NO:256) M13903 CDS 3.43 679 AGCAGCAGGAGGGGAGCTGGAGCTCTCTG (SEQ ID NO:257) M14058 3'UTR 12 2221 A M14083 3'UTR 7 2667 AT M14170 CDS 7.66 30 TGC M14219 3'UTR 13 1660 T M14630 UNKNOWN 12 538 A M14648 3'UTR 10.66 3535 TTG M14745 3'UTR 7.5 897 AC M14764 5'UTR 3.6 36 AGCGC M14764 3'UTR 7 2444 CA M15169 UNKNOWN 13 2852 C M15353 3'UTR 18 847 T M16276 UNKNOWN 12 1368 A M16505 UNKNOWN 6.5 2611 AC M16505 UNKNOWN 18 3645 A M16801 CDS 3.83 2295 CCCCCA M16937 3'UTR 2.7 865 AAACAAA (SEQ ID NO:258) M16938 5'UTR 7 172 TG M16965 CDS 2.94 115 TACCTTTGTTGGAAGACG (SEQ ID NO:259) M16965 CDS 2.5 591 CTGGAAGACATGGATTTT (SEQ ID NO:260) M18533 UNKNOWN 5.25 12297 TTGA M18533 UNKNOWN 8.5 11725 AC M18728 UNKNOWN 12 2266 A M19154 3'UTR 7.33 2117 ACA M19961 5'UTR 13 0 T M20681 UNKNOWN 13 2233 T M20681 UNKNOWN 13 3705 A M21305 CDS 6.4 56 TGGAA M21305 BORDER 3.8 0 ATGGA M21574 3'UTR 14 4503 T M23114 3'UTR 3.8 3839 CACCC M23263 CDS 20.33 1884 GGC M23263 CDS 17 701 GCA M24069 CDS 7.66 229 CCA M24283 3'UTR 9 2742 GT M24486 3'UTR 16 2350 T M24902 3'UTR 12.5 2338 AATA M25667 3'UTR 9 1153 CT M25667 3'UTR 17 973 A M28170 3'UTR 17 1816 GT M28713 5'UTR 6.66 253 CGG M29053 3'UTR 3.8 1579 AAAAT M29204 UNKNOWN 21 288 A M29873 3'UTR 6.5 1687 TA M29874 3'UTR 7 1688 AT M29874 3'UTR 15 2510 T M30448 3'UTR 26 887 A M31165 BORDER 12 900 A M31523 3'UTR 12 2316 T M31525 3'UTR 6.5 1037 AC M31682 3'UTR 8 1601 AG M31682 3'UTR 6.5 1886 TG M31732 3'UTR 15 1474 C M31899 CDS 5.66 869 GAA M31932 3'UTR 13 1430 T M32315 3'UTR 7.75 1814 TTTG M32315 3'UTR 6.5 3589 TG M32315 3'UTR 12 2368 A M34041 CDS 6.66 903 GAG M34309 5'UTR 6.5 4 CA M34539 3'UTR 12 1120 T M35531 3'UTR 6.25 2068 TTAT M35531 3'UTR 18 1752 T M35663 3'UTR 18 2056 T M36089 UNKNOWN 17 2462 AC M36542 3'UTR 18 1886 A M36711 3'UTR 6.66 1494 GCC M36820 3'UTR 6.25 540 TATT M36860 CDS 3.66 950 CAGCTG M37981 CDS 6.33 113 GCT M54915 5'UTR 3.8 246 CAGCA M54915 5'UTR 9 45 GCA M54915 3'UTR 5.25 2191 TATT M54927 3'UTR 14 2476 A M55047 3'UTR 22.5 2095 GT M55053 3'UTR 12 1915 T M55172 CDS 7.96 3238 CTGCCCCCTGGAGTAGAGGACATCAGCGGGCTTCCTTCTG GAGAAGTTCTAGAGACCG (SEQ ID NO:261) M55172 CDS 5.52 2979 GGGCTTCCTTCTGGAGAAGTTCTAGAGACCACTGCCCCCT GGAGTAGAGGACATCAGC (SEQ ID NO:262) M55172 CDS 5 3636 GCTGCCCCCTGGAGTAGAGGACATCAGCGGGCTTCCTTCT GGAGAAGTTCTAGAGACT (SEQ ID NO:263) M55422 5'UTR 5.66 944 TTG M55542 3'UTR 14 2006 A M55593 5'UTR 7.33 87 GCG M55630 UNKNOWN 22 126 GT M55630 UNKNOWN 12 1538 A M55654 CDS 18.66 466 CAG M55654 CDS 9.66 430 CAG M55654 3'UTR 13 1297 T M55683 3'UTR 12 1203 TG M57627 3'UTR 3.5 1457 AAAAAT M58583 3'UTR 5.66 1354 TTG M59305 5'UTR 4.16 129 CTTTTT M59465 3'UTR 12 3986 A M59499 3'UTR 2.91 2093

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L5: Entry 11 of 16

File: USPT

Apr 10, 2001

DOCUMENT-IDENTIFIER: US 6214795 B1

TITLE: Peptide compounds useful for modulating FGF receptor activity

DATE FILED (1):19961112Brief Summary Text (4):

The FGFs mediate their effects by binding to high affinity cell surface receptors (reviewed in Johnson and Williams (1992) Adv. Cancer Res. 60:1-41). Four distinct FGF receptors have been identified: FGFR1 (also known as Flg, bFGFR, Cek1 or N-bFGFR) (Lee et al. (1989) Science 245:57-60; Dionne et al. (1990) EMBO J. 9:2685-2692; Johnson et al. (1990) Mol. Cell. Biol. 10:4728-4736; Eisemann et al. (1991) Oncogene 6:1195-1202; Hou et al. (1991) Science 251:665-668), FGFR2 (also known as Bek, Cek3, K-sam, TK14, TK25 or KGFR) (Dionne et al. (1990) EMBO J. 9:2685-2692; Hattori et al. (1990) Proc. Natl. Acad. Sci. USA 87:5983-5987; Miki et al. (1991) Science 251:72-75; Saiki et al. (1988) Science 239:487-491; Pasquale (1990) Proc. Natl. Acad. Sci. USA 87:5812-5816; Houssaint et al. (1990) Proc. Natl. Acad. Sci. USA 87:8180-8184; Champion-Arnaud et al. (1991) Oncogene 6:979-987; Crumley et al. (1991) Oncogene 6:2255-2262; Raz et al. (1991) Oncogene 6:753-760; Sato et al. (1991) Oncogene 6:1279-1283), FGFR3 (also known as Cek2) (Keegan et al. (1991) Proc. Natl. Acad. Sci. USA 88:1095-1099) and FGFR4 (Partanen et al. (1991) EMBO J. 10:1347-1354).

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Proc Natl Acad Sci U S A 1991 Feb 15;88(4):1095-9

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Isolation of an additional member of the fibroblast growth factor receptor family, FGFR-3.

Keegan K, Johnson DE, Williams LT, Hayman MJ.

Department of Microbiology, State University of New York, Stony Brook 11794.

The fibroblast growth factors are a family of polypeptide growth factors involved in a variety of activities including mitogenesis, angiogenesis, and wound healing.

Fibroblast growth factor receptors (FGFRs) have previously been identified in chicken, mouse, and human and have been shown to contain an extracellular domain with either two or three immunoglobulin-like domains, a transmembrane domain, and a cytoplasmic tyrosine kinase domain. We have isolated a human cDNA for another tyrosine kinase receptor that is highly homologous to the previously described FGFR. Expression of this receptor cDNA in COS cells directs the expression of a 125-kDa glycoprotein. We demonstrate that this cDNA encodes a biologically active receptor by showing that human acidic and basic fibroblast growth factors activate this receptor as measured by  $^{45}\text{Ca}^{2+}$  efflux assays. These data establish the existence of an additional member of the FGFR family that we have named FGFR-3.

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File: USPT

Jul 21, 1998

DOCUMENT-IDENTIFIER: US 5783683 A

TITLE: Antisense oligonucleotides which reduce expression of the FGFR1 gene

DATE FILED (1):19950110Drawing Description Text (10):

FIG. 9 shows an RT-PCR Southern Blot of FGFR1, FGFR3, and FGFR4 demonstrating the selective reduction of FGFR1 mRNA following treatment of glioblastoma cells with the antisense molecules of the invention.

Detailed Description Text (122):

mRNA was analyzed as described above with the exception that both FGFR1, FGFR3 and FGFR4 mRNA were studied in this particular work. SNB-19 glioblastoma cells were plated at 1.times.10.sup.5 cells per 100 mm dish in serum-supplemented medium. Eighteen hours later the cells were converted to serum-free medium containing FGFR1.alpha. antisense oligonucleotide (R1AS.alpha., 30 .mu.m) or FGFR1.alpha. antisense reverse control oligonucleotide (R1.alpha.RC, 30 .mu.m). Non-treated cells (NT) were run as a control. Cells were treated for three consecutive days with oligonucleotide. Cells were scraped on day 7 and mRNA and cDNA were purified and synthesized respectively. Using cDNA from each of the three different treatments, PCR was used to amplify cDNA for FGFR1, FGFR3, and FGFR4 receptors. SNB-19 cells do not produce FGFR2.





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**Targeted expression of a human pituitary tumor-derived isoform of FGF receptor-4 recapitulates pituitary tumorigenesis.**

Ezzat S, Zheng L, Zhu XF, Wu GE, Asa SL.

*J Clin Invest.* 2002 Jan;109(1):69-78.

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**Histological and genetic diagnosis of gliomatosis cerebri: case report.**

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*Anal Quant Cytol Histol.* 2000 Jun;22(3):267-74.

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**Suppression of glioblastoma cell growth following antisense oligonucleotide-mediated inhibition of fibroblast growth factor receptor expression.**

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*Glia.* 1999 Oct;28(1):66-76.

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- 21 ☐ **Basic fibroblast growth factor and fibroblast growth factor receptor implicated in the growth of human astrocytomas.**  
Morrison RS, Yamaguchi F, Saya H, Bruner JM, Yahanda AM, Donehow LA, Berger M.  
*J Neurooncol.* 1994;18(3):207-16.  
PMID: 7964981 [PubMed - indexed for MEDLINE]  
From PubMed
- 22 ☐ **Identification and characterization of high molecular weight forms of basic fibroblast growth factor in human pituitary adenomas.**  
Li Y, Koga M, Kasayama S, Matsumoto K, Arita N, Hayakawa T, Sato B  
*J Clin Endocrinol Metab.* 1992 Dec;75(6):1436-41.  
PMID: 1464644 [PubMed - indexed for MEDLINE]  
From PubMed
- 23 ☐ **Angiogenic activity of the K-fgf/hst oncogene in neural transplants.**  
Brustle O, Aguzzi A, Talarico D, Basilico C, Kleihues P, Wiestler OD.  
*Oncogene.* 1992 Jun;7(6):1177-83.  
PMID: 1375717 [PubMed - indexed for MEDLINE]  
From PubMed
- 24 ☐ **Phosphorothioate antisense oligonucleotides against basic fibroblast growth factor inhibit anchorage-dependent and anchorage-independent growth of a malignant glioblastoma cell line.**  
Murphy PR, Sato Y, Knee RS.  
*Mol Endocrinol.* 1992 Jun;6(6):877-84.  
PMID: 1323055 [PubMed - indexed for MEDLINE]  
From PubMed
- 25 ☐ **Gene expression of fibroblast growth factor receptors in the tissues of human gliomas and meningiomas.**  
Takahashi JA, Suzui H, Yasuda Y, Ito N, Ohta M, Jaye M, Fukumoto M, Y, Kikuchi H, Hatanaka M.  
*Biochem Biophys Res Commun.* 1991 May 31;177(1):1-7.  
PMID: 1645953 [PubMed - indexed for MEDLINE]  
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M64347. Human novel growt...[gi:182564]

## Links

LOCUS HUMFGFLR 3829 bp mRNA linear PRI 31-DEC-1994  
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 VERSION M64347.1 GI:182564  
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 ORGANISM Homo sapiens  
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 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 REFERENCE 1 (bases 1 to 3829)  
 AUTHORS Thompson,L.M., Plummer,S., Schalling,M., Altherr,M.R.,  
 Gusella,J.F., Housman,D.E. and Wasmuth,J.J.  
 TITLE A gene encoding a fibroblast growth factor receptor isolated from  
 the Huntington disease gene region of human chromosome 4  
 JOURNAL Genomics 11 (4), 1133-1142 (1991)  
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 PUBMED 1664411  
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NM\_000142. Homo sapiens fibr...[gi:13112046]

# Links

LOCUS FGFR3 4093 bp mRNA linear PRI 21-FEB-2001

DEFINITION Homo sapiens fibroblast growth factor receptor 3 (achondroplasia, thanatophoric dwarfism) (FGFR3), transcript variant 1, mRNA.

ACCESSION NM\_000142

VERSION NM\_000142.2 GI:13112046

KEYWORDS .

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 4093)

AUTHORS Partanen,J., Makela,T.P., Alitalo,R., Lehvaslaiho,H. and Alitalo,K.

TITLE Putative tyrosine kinases expressed in K-562 human leukemia cells

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 87 (22), 8913-8917 (1990)

MEDLINE 91062389

PUBMED 2247464

REFERENCE 2 (bases 1 to 4093)

AUTHORS Keegan,K., Johnson,D.E., Williams,L.T. and Hayman,M.J.

TITLE Isolation of an additional member of the fibroblast growth factor receptor family, FGFR-3

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 88 (4), 1095-1099 (1991)

MEDLINE 91142118

PUBMED 1847508

REFERENCE 3 (bases 1 to 4093)

AUTHORS Thompson,L.M., Plummer,S., Schalling,M., Altherr,M.R., Gusella,J.F., Housman,D.E. and Wasmuth,J.J.

TITLE A gene encoding a fibroblast growth factor receptor isolated from the Huntington disease gene region of human chromosome 4

JOURNAL Genomics 11 (4), 1133-1142 (1991)

MEDLINE 92147110

PUBMED 1664411

REFERENCE 4 (bases 1 to 4093)

AUTHORS Velinov,M., Slaugenhaupt,S.A., Stoilov,I., Scott,C.I. Jr., Gusella,J.F. and Tsipouras,P.

TITLE The gene for achondroplasia maps to the telomeric region of chromosome 4p

JOURNAL Nat. Genet. 6 (3), 314-317 (1994)

MEDLINE 94282083

PUBMED 8012397

REFERENCE 5 (bases 1 to 4093)

AUTHORS Le Merrer,M., Rousseau,F., Legeai-Mallet,L., Landais,J.C., Pelet,A., Bonaventure,J., Sanak,M., Weissenbach,J., Stoll,C., Munnich,A. et al.

TITLE A gene for achondroplasia-hypochondroplasia maps to chromosome 4p

JOURNAL Nat. Genet. 6 (3), 318-321 (1994)

MEDLINE 94282084  
PUBMED 8012398  
REFERENCE 6 (bases 1 to 4093)  
AUTHORS Francomano,C.A., Ortiz de Luna,R.I., Hefferon,T.W., Bellus,G.A.,  
Turner,C.E., Taylor,E., Meyers,D.A., Blanton,S.H., Murray,J.C.,  
McIntosh,I. et al.  
TITLE Localization of the achondroplasia gene to the distal 2.5 Mb of  
human chromosome 4p  
JOURNAL Hum. Mol. Genet. 3 (5), 787-792 (1994)  
MEDLINE 94362678  
PUBMED 8081365  
REFERENCE 7 (bases 1 to 4093)  
AUTHORS Perez-Castro,A.V., Wilson,J. and Altherr,M.R.  
TITLE Genomic organization of the human fibroblast growth factor receptor  
3 (FGFR3) gene and comparative sequence analysis with the mouse  
Fgfr3 gene  
JOURNAL Genomics 41 (1), 10-16 (1997)  
MEDLINE 97271550  
PUBMED 9126476  
REFERENCE 8 (bases 1 to 4093)  
AUTHORS Passos-Bueno,M.R., Wilcox,W.R., Jabs,E.W., Sertie,A.L., Alonso,L.G.  
and Kitoh,H.  
TITLE Clinical spectrum of fibroblast growth factor receptor mutations  
JOURNAL Hum. Mutat. 14 (2), 115-125 (1999)  
MEDLINE 99355711  
PUBMED 10425034  
COMMENT REVIEWED REFSEQ: This record has been curated by NCBI staff. The  
reference sequence was derived from M58051.1 and M64347.1.  
On Feb 23, 2001 this sequence version replaced gi:4503710.  
Summary: The protein encoded by this gene is a member of the  
fibroblast growth factor receptor family, where amino acid sequence  
is highly conserved between members and throughout evolution. FGFR  
family members differ from one another in their ligand affinities  
and tissue distribution. A full-length representative protein would  
consist of an extracellular region, composed of three  
immunoglobulin-like domains, a single hydrophobic membrane-spanning  
segment and a cytoplasmic tyrosine kinase domain. The extracellular  
portion of the protein interacts with fibroblast growth factors,  
setting in motion a cascade of downstream signals, ultimately  
influencing mitogenesis and differentiation. This particular family  
member binds acidic and basic fibroblast growth hormone and plays a  
role in bone development and maintenance. Mutations in this gene  
lead to craniosynostosis and multiple types of skeletal dysplasia.  
Alternative splicing occurs and additional variants have been  
described, including those utilizing alternate exon 8 rather than  
9, but their full-length nature has not been determined.  
Transcript Variant: This variant (1) is missing alternatively  
spliced exon 8 but utilizes alternatively spliced exon 9, resulting  
in isoform (1) with the IIIc-type C-terminal half of the IgIII  
domain.



COMPLETENESS: complete on the 3' end.

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NM\_022965. Homo sapiens fibr...[gi:13112047]

## Links

LOCUS FGFR3 3757 bp mRNA linear PRI 21-FEB-2001

DEFINITION Homo sapiens fibroblast growth factor receptor 3 (achondroplasia, thanatophoric dwarfism) (FGFR3), transcript variant 2, mRNA.

ACCESSION NM\_022965

VERSION NM\_022965.1 GI:13112047

KEYWORDS .

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 3757)

AUTHORS Partanen,J., Makela,T.P., Alitalo,R., Lehvaslaiho,H. and Alitalo,K.

TITLE Putative tyrosine kinases expressed in K-562 human leukemia cells

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 87 (22), 8913-8917 (1990)

MEDLINE 91062389

PUBMED 2247464

REFERENCE 2 (bases 1 to 3757)

AUTHORS Keegan,K., Johnson,D.E., Williams,L.T. and Hayman,M.J.

TITLE Isolation of an additional member of the fibroblast growth factor receptor family, FGFR-3

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 88 (4), 1095-1099 (1991)

MEDLINE 91142118

PUBMED 1847508

REFERENCE 3 (bases 1 to 3757)

AUTHORS Thompson,L.M., Plummer,S., Schalling,M., Altherr,M.R., Gusella,J.F., Housman,D.E. and Wasmuth,J.J.

TITLE A gene encoding a fibroblast growth factor receptor isolated from the Huntington disease gene region of human chromosome 4

JOURNAL Genomics 11 (4), 1133-1142 (1991)

MEDLINE 92147110

PUBMED 1664411

REFERENCE 4 (bases 1 to 3757)

AUTHORS Velinov,M., Slaugenhaupt,S.A., Stoilov,I., Scott,C.I. Jr., Gusella,J.F. and Tsipouras,P.

TITLE The gene for achondroplasia maps to the telomeric region of chromosome 4p

JOURNAL Nat. Genet. 6 (3), 314-317 (1994)

MEDLINE 94282083

PUBMED 8012397

REFERENCE 5 (bases 1 to 3757)

AUTHORS Le Merrer,M., Rousseau,F., Legeai-Mallet,L., Landais,J.C., Pelet,A., Bonaventure,J., Sanak,M., Weissenbach,J., Stoll,C., Munnich,A. et al.

TITLE A gene for achondroplasia-hypochondroplasia maps to chromosome 4p

JOURNAL Nat. Genet. 6 (3), 318-321 (1994)

MEDLINE 94282084  
 PUBMED 8012398  
 REFERENCE 6 (bases 1 to 3757)  
 AUTHORS Francomano,C.A., Ortiz de Luna,R.I., Hefferon,T.W., Bellus,G.A.,  
 Turner,C.E., Taylor,E., Meyers,D.A., Blanton,S.H., Murray,J.C.,  
 McIntosh,I. et al.  
 TITLE Localization of the achondroplasia gene to the distal 2.5 Mb of  
 human chromosome 4p  
 JOURNAL Hum. Mol. Genet. 3 (5), 787-792 (1994)  
 MEDLINE 94362678  
 PUBMED 8081365  
 REFERENCE 7 (bases 1 to 3757)  
 AUTHORS Perez-Castro,A.V., Wilson,J. and Altherr,M.R.  
 TITLE Genomic organization of the human fibroblast growth factor receptor  
 3 (FGFR3) gene and comparative sequence analysis with the mouse  
 Fgfr3 gene  
 JOURNAL Genomics 41 (1), 10-16 (1997)  
 MEDLINE 97271550  
 PUBMED 9126476  
 REFERENCE 8 (bases 1 to 3757)  
 AUTHORS Passos-Bueno,M.R., Wilcox,W.R., Jabs,E.W., Sertie,A.L., Alonso,L.G.  
 and Kitoh,H.  
 TITLE Clinical spectrum of fibroblast growth factor receptor mutations  
 JOURNAL Hum. Mutat. 14 (2), 115-125 (1999)  
 MEDLINE 99355711  
 PUBMED 10425034  
 COMMENT REVIEWED REFSEQ: This record has been curated by NCBI staff. The  
 reference sequence was derived from AF245114.1 and M64347.1.  
 Summary: The protein encoded by this gene is a member of the  
 fibroblast growth factor receptor family, where amino acid sequence  
 is highly conserved between members and throughout evolution. FGFR  
 family members differ from one another in their ligand affinities  
 and tissue distribution. A full-length representative protein would  
 consist of an extracellular region, composed of three  
 immunoglobulin-like domains, a single hydrophobic membrane-spanning  
 segment and a cytoplasmic tyrosine kinase domain. The extracellular  
 portion of the protein interacts with fibroblast growth factors,  
 setting in motion a cascade of downstream signals, ultimately  
 influencing mitogenesis and differentiation. This particular family  
 member binds acidic and basic fibroblast growth hormone and plays a  
 role in bone development and maintenance. Mutations in this gene  
 lead to craniosynostosis and multiple types of skeletal dysplasia.  
 Alternative splicing occurs and additional variants have been  
 described, including those utilizing alternate exon 8 rather than  
 9, but their full-length nature has not been determined.  
 Transcript Variant: This variant (2) does not contain alternatively  
 spliced exons 8 or 9, resulting in a loss of the C-terminal half of  
 the IgIII domain. In addition, this variant is missing  
 alternatively spliced exon 10 which encodes the transmembrane  
 region, suggesting a soluble receptor.

COMPLETENESS: complete on the 3' end.

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gi|13112047|ref|NM\_022965.1| Homo sapiens fibroblast growth factor receptor 3  
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thanatophoric dwarfism) (FGFR3), transcript variant 2,  
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M58051. Human fibroblast ...[gi:182568]

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 VERSION M58051.1 GI:182568  
 KEYWORDS fibroblast growth factor receptor.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 REFERENCE 1 (bases 1 to 2520)  
 AUTHORS Keegan,K., Johnson,D.E., Williams,L.T. and Hayman,M.J.  
 TITLE Isolation of an additional member of the fibroblast growth factor  
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 JOURNAL Proc. Natl. Acad. Sci. U.S.A. 88 (4), 1095-1099 (1991)  
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 PUBMED 1847508  
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mRNA, complete cds

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Identities = 2244/2256 (99%)

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AF487554. Homo sapiens fibr...[gi:20452379]

## Links

LOCUS AF487554 16976 bp DNA linear PRI 05-MAY-2002

DEFINITION Homo sapiens fibroblast growth factor receptor 3 (FGFR3) gene, partial cds, alternatively spliced.

ACCESSION AF487554

VERSION AF487554.1 GI:20452379

KEYWORDS .

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 16976)

AUTHORS Lind,D.L. and Cox,D.R.

TITLE Fibroblast growth factor receptor 3 (FGFR3) genomic sequence

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 16976)

AUTHORS Lind,D.L. and Cox,D.R.

TITLE Direct Submission

JOURNAL Submitted (26-FEB-2002) Genetics, Stanford University, SUMC M312, 300 Pasteur Drive M/C 5120, Stanford, CA 94305-5120, USA

FEATURES Location/Qualifiers

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F245114. Homo sapiens fibr...[gi:7533124]

## Links

LOCUS AF245114 2184 bp mRNA linear PRI 28-MAR-2002

DEFINITION Homo sapiens fibroblast growth factor receptor 3 (FGFR3) mRNA, complete cds, alternatively spliced.

ACCESSION AF245114

VERSION AF245114.1 GI:7533124

KEYWORDS .

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 2184)

AUTHORS Terada,M., Shimizu,A., Sato,N., Miyakaze,S.I., Katayama,H. and Kurokawa-Seo,M.

TITLE Fibroblast growth factor receptor 3 lacking the Ig IIIb and transmembrane domains secreted from human squamous cell carcinoma DJM-1 binds to FGFs

JOURNAL Mol. Cell Biol. Res. Commun. 4 (6), 365-373 (2001)

MEDLINE 21561228

PUBMED 11703096

REFERENCE 2 (bases 1 to 2184)

AUTHORS Terada,M., Shimizu,A. and Seo,M.

TITLE Secretion and dimerization of the FGFR3 isoform, resulting from alternative splicing, that is expressed in human malignant trichilemmal cyst cell

JOURNAL Unpublished

REFERENCE 3 (bases 1 to 2184)

AUTHORS Terada,M., Shimizu,A. and Seo,M.

TITLE Direct Submission

JOURNAL Submitted (14-MAR-2000) Biotechnology, Kyoto Sangyo University, Kamigamo-Motoyama, Kita-Ku, Kyoto 603-8555, Japan

FEATURES

Location/Qualifiers

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OR  
L5 132127 S (CEREBRAL OR (CHOROID(W)PLEXUS) OR CEREBELLUM OR  
HYPOTHALAMIC  
L6 10081 S (CEREBRAL OR (CHOROID(W)PLEXUS) OR CEREBELLUM OR  
HYPOTHALAMIC  
L7 204265 S L4 OR L5 OR L6  
L8 151 S L3 AND L7  
L9 115 S L8 AND PY<2001  
L10 136 S L8 AND PY<2002  
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L12 1822 S (FGFR3 OR (FGF(W)R3) OR (FGF(W)RECEPTOR(W)3) OR  
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L13 15 S L12 AND L7  
L14 6 DUP REM L13 (9 DUPLICATES REMOVED)  
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L16 0 S L15 AND L8  
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10/ 066,305

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DOCUMENT NUMBER: 20349276 PubMed ID: 10889045  
TITLE: Repeat polymorphisms within gene regions: phenotypic and evolutionary implications.  
AUTHOR: Wren J D; Forgacs E; Fondon J W 3rd; Pertsemlidis A; Cheng S Y; Gallardo T; Williams R S; Shohet R V; Minna J D; Garner H R  
CORPORATE SOURCE: Program in Genetics, Southwestern Graduate School of Biomedical Sciences, Dallas, TX, USA.  
CONTRACT NUMBER: P50CA70907 (NCI)  
SOURCE: AMERICAN JOURNAL OF HUMAN GENETICS, (2000 Aug) 67 (2) 345-56.  
Journal code: 0370475. ISSN: 0002-9297.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
OTHER SOURCE: GENBANK-AF013956; GENBANK-AF017789; GENBANK-AF032886; GENBANK-AF042838; GENBANK-AF047437; GENBANK-AF060231; GENBANK-D14838; GENBANK-D83492; GENBANK-D86407; GENBANK-D86550; GENBANK-L08835; GENBANK-M55047; GENBANK-M60052; GENBANK-M60315; GENBANK-M64347; GENBANK-R12160; GENBANK-R42196; GENBANK-T47177; GENBANK-T62484; GENBANK-T63962; GENBANK-T70173; GENBANK-U49020; GENBANK-U60325; GENBANK-X55313; GENBANK-X70811; GENBANK-X78261; GENBANK-X82209; GENBANK-Y00285; GENBANK-Y11525; +  
ENTRY MONTH: 200008  
ENTRY DATE: Entered STN: 20000901  
Last Updated on STN: 20030105  
Entered Medline: 20000821

AB We have developed an algorithm that predicted 11,265 potentially polymorphic tandem repeats within transcribed sequences. We estimate that 22% (2,207/9,717) of the annotated clusters within UniGene contain at least one potentially polymorphic locus. Our predictions were tested by allelotyping a panel of approximately 30 individuals for 5% of these regions, confirming polymorphism for more than half the loci tested. Our study indicates that tandem-repeat polymorphisms in genes are more common than is generally believed. Approximately 8% of these loci are within coding sequences and, if polymorphic, would result in frameshifts. Our catalogue of putative polymorphic repeats within transcribed sequences comprises a large set of potentially phenotypic or disease-causing loci. In addition, from the anomalous character of the repetitive sequences within unannotated clusters, we also conclude that the UniGene cluster count substantially overestimates the number of genes in the human genome.

We hypothesize that polymorphisms in repeated sequences occur with some baseline distribution, on the basis of repeat homogeneity, size, and sequence composition, and that deviations from that distribution are indicative of the nature of selection pressure at that locus. We find evidence of selective maintenance of the ability of some genes to respond very rapidly, perhaps even on intragenerational timescales, to fluctuating selective pressures.

L2 ANSWER 4 OF 7 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2001:758742 CAPLUS

DOCUMENT NUMBER: 135:314390

TITLE: Large-scale monitoring of expression patterns of p53-regulated gene and analysis of p53 gene function

INVENTOR(S): Mack, David H.

PATENT ASSIGNEE(S): Affymetrix, Inc., USA

SOURCE: U.S., 46 pp., Cont.-in-part of Appl. No. PCT/US98/01206.

CODEN: USXXAM

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6303301	B1	20011016	US 1998-86285	19980529
WO 9830722	A1	19980716	WO 1998-US1206	19980112

W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG

US 2002028454	A1	20020307	US 2001-836278	20010418
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PRIORITY APPLN. INFO.: US 1997-35327P P 19970113  
WO 1998-US1206 A2 19980112  
US 1997-49627P P 19970613  
US 1998-86285 A3 19980529

AB This invention provides methods, compns. and app. for mapping the regulatory relationships of genes by massive parallel monitoring of gene expression. The information obtained can be of use in drug discovery (no data). The method uses high d. oligonucleotide arrays to monitor changes in expression in response to events and stimuli. Very large nos. of gene (>6,500) may be monitored in this method using samples from many tissues and developmental or disease stages. Changes are quantified and a relationship model constructed using LISREL (Linear Structure Relationship) methods. Mutations in up-stream regulatory genes can be detected by monitoring the change in down-stream gene expression. Similarly, the effect of a specific mutation in an up-stream gene is detd. by monitoring the down-stream gene expression. In addn., regulatory function of a target gene can be detd. by monitoring the expression of a large no. of down-stream genes. The invention also provides specific embodiments for detecting p53 functional homozygous and heterozygous mutations and for detg. the function of p53 mutations.

REFERENCE COUNT: 46 THERE ARE 46 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE REFORMAT

L2 ANSWER 1 OF 7 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2002:241013 CAPLUS

DOCUMENT NUMBER: 136:277466

TITLE: Expressed gene sets as markers for specific tumors

INVENTOR(S): Ramaswamy, Sridhar; Golub, Todd B.; Tamayo, Pablo; Angelo, Michael

PATENT ASSIGNEE(S): Whitehead Institute for Biomedical Research, USA; Dana-Farber Cancer Institute, Inc.

SOURCE: PCT Int. Appl., 715 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 4

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002024956	A2	20020328	WO 2001-US29287	20010919
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
WO 2002024956	A2	20020328	WO 2001-XA29287	20010919
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
WO 2002024956	A2	20020328	WO 2001-XB29287	20010919
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
WO 2002024956	A2	20020328	WO 2001-XC29287	20010919
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			

AU 2001092802	A5	20020402	AU 2001-92802	20010919
US 2002110820	A1	20020815	US 2001-955920	20010919
PRIORITY APPLN. INFO.:			US 2000-233534P	P 20000919
			US 2001-278749P	P 20010326
			WO 2001-US29287	W 20010919

AB Sets of genetic markers for specific tumor classes are described, as well as methods of identifying a biol. sample based on these markers. Total RNA was isolated from .apprx.300 human tumor and normal tissue specimens representing 30 individual classes of tumor or normal tissue, and cDNA produced using established mol. biol. protocols was hybridized to two

high d. Affymetrix oligonucleotide microarrays (Hu6800FL and Hu35KsubA0). Raw expression data was combined into a master data set contg. the expression values for between 6800 and 16,000 genes expressed by each individual sample. A filter was applied to this data set which only allows those genes expressed at 3-fold above baseline and with an abs. difference in expression value of 100 to pass. By comparing the sets of genes which

are expressed specifically in one class of tumor (e.g., pancreatic adenocarcinoma) vs. its accompanying normal tissue (e.g., normal pancreas), sets of genes were detd. which are specific to various tumors and their normal tissue counterparts. Also described are diagnostic, prognostic, and therapeutic screening uses for these markers, as well as oligonucleotide arrays comprising these markers. [This abstr. record is one of 4 records for this document necessitated by the large no. of index entries required to fully index the document and publication system constraints.].

L2 ANSWER 2 OF 7 CAPLUS COPYRIGHT 2003 ACS



L2 ANSWER 7 OF 7

MEDLINE

DUPLICATE 1

ACCESSION NUMBER: 92147110 MEDLINE  
DOCUMENT NUMBER: 92147110 PubMed ID: 1664411  
TITLE: A gene encoding a fibroblast growth factor receptor  
isolated from the Huntington disease gene region of human  
chromosome 4.  
AUTHOR: Thompson L M; Plummer S; Schalling M; Altherr M R; Gusella  
J F; Housman D E; Wasmuth J J  
CORPORATE SOURCE: Department of Biological Chemistry, College of Medicine,  
University of California, Irvine 92717.  
CONTRACT NUMBER: NS25631-04 (NINDS)  
SOURCE: GENOMICS, (1991 Dec) 11 (4) 1133-42.  
Journal code: 8800135. ISSN: 0888-7543.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
OTHER SOURCE: GENBANK-M64347  
ENTRY MONTH: 199203  
ENTRY DATE: Entered STN: 19920405  
Last Updated on STN: 20000303  
Entered Medline: 19920313

AB The gene responsible for Huntington disease (HD), an autosomal dominant  
neurodegenerative disorder, is located near the terminus of the short arm  
of chromosome 4. Detailed genetic linkage and physical mapping studies  
have defined a region of approximately 2.5 million basepairs where the  
disease gene is likely to be located. Efforts to identify the disease  
gene  
are now focused on the identification and characterization of expressed  
genes in this region. Nucleotide sequence analysis of a cDNA clone  
derived  
from the HD gene region has revealed that it encodes a member of the  
fibroblast growth factor subfamily of tyrosine kinase receptors, some  
members of which are known to be involved in the differentiation and  
survival of certain cell types within the central nervous system.  
Histochemical analysis using in situ hybridization revealed its  
expression  
in many areas of the brain, among them being the caudate and putamen. The  
nature of this gene, FGFR3, and its map location make it a possible  
candidate for the HD gene.

L2 ANSWER 6 OF 7 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1999:641026 CAPLUS  
DOCUMENT NUMBER: 131:267987  
TITLE: Cancer diagnosis and therapy based on expression levels of p53-regulated genes  
INVENTOR(S): Levine, Arnold J.; Murphy, Maureen E.; Mack, David H.;  
Gish, Kurt C.; Tom, Edward Yat Wah  
PATENT ASSIGNEE(S): Affymetrix, Inc., USA; Princeton University  
SOURCE: PCT Int. Appl., 46 pp.  
CODEN: PIXXD2  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9950456	A1	19991007	WO 1999-US6656	19990326
W:				
AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW:				
GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
US 6020135	A	20000201	US 1998-49025	19980327
CA 2324444	AA	19991007	CA 1999-2324444	19990326
AU 9932085	A1	19991018	AU 1999-32085	19990326
EP 1064404	A1	20010103	EP 1999-914184	19990326
R:				
AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				
US 6171798	B1	20010109	US 1999-442039	19991117
US 2001039013	A1	20011108	US 2001-755028	20010108
PRIORITY APPLN. INFO.:			US 1998-49025	A1 19980327
			WO 1999-US6656	W 19990326
			US 1999-442039	A3 19991117

AB Many genes are identified as being p53-regulated which were not heretofore

known to be p53-regulated. This includes both genes whose expression is induced and genes whose expression is repressed by the expression of wild-type p53. The effects of p53 expression on gene expression in Eb-1 cells was tested by hybridizing to a chip that contains deoxyoligonucleotide sequences (25-mers) that derived from a database of 6800 known genes or EST sequences. Seventy genes were induced by p53 and 77 were repressed by p53. Monitoring expression of these genes is used

to provide indications of p53 status in a cell. Such monitoring can also be used to screen for useful anticancer therapeutics, as well as for substances which are carcinogenic. Defects in p53 can be bypassed by supplying p53 induced genes to cells. Defects in p53 can also be

bypassed by supplying antisense constructs to p53-repressed genes.

REFERENCE COUNT: 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

L11 ANSWER 66 OF 67 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1991:79455 CAPLUS

DOCUMENT NUMBER: 114:79455

TITLE: Suppression of basic fibroblast growth factor  
expression by antisense oligodeoxynucleotides

inhibits

the growth of transformed human astrocytes

AUTHOR(S): Morrison, Richard S.

CORPORATE SOURCE: Robert S. Dow Neurol. Sci. Inst., Good Samaritan  
Hosp., Portland, OR, 97209, USA

SOURCE: Journal of Biological Chemistry (1991),  
266(2), 728-34

CODEN: JBCHA3; ISSN: 0021-9258

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Basic fibroblast growth factor (bFGF) is a heparin-binding protein  
expressing potent mitogenic and angiogenic properties. Elevated levels  
of

bFGF have recently been described in human **glioma** cell lines.

The high degree of vascularity and invasiveness which characterize human  
**gliomas** suggest that activated expression of bFGF or similar  
proteins may be related to the aberrant growth patterns of these tumors.  
The influence of endogenous bFGF on **glioma** cell growth in vitro  
was evaluated in the present study by downregulating bFGF expression

using

antisense oligonucleotide primers. The addn. of 50 .mu.M bFGF-specific  
antisense primer to the human **glioma** cell line SNB-19 resulted  
in an 80% inhibition in **glioma** growth. This effect was  
saturable and specific. Antisense primers directed to 2 different sites  
of bFGF mRNA were effective in suppressing SNB-19 growth, whereas sense  
strand primer was ineffective. Furthermore, only the antisense primer  
significantly reduced the specific activity of bFGF protein in SNB-19

cell

exts. Neither antisense or sense primers inhibited the growth of  
non-transformed human glia. BFGF mRNA was detected in both transformed  
and nontransformed human glia by polymerase chain reaction anal.  
suggesting that alterations in bFGF isoform content or activity may be  
specifically related to abnormal growth control in human **gliomas**

L11 ANSWER 65 OF 67

MEDLINE

DUPLICATE 26

ACCESSION NUMBER: 91342665 MEDLINE

DOCUMENT NUMBER: 91342665 PubMed ID: 1652059

TITLE: The human **fibroblast growth factor receptor** genes: a common structural arrangement underlies the mechanisms for generating receptor forms that differ in their third immunoglobulin domain.

AUTHOR: Johnson D E; Lu J; Chen H; Werner S; Williams L T

CORPORATE SOURCE: Howard Hughes Medical Institute, Program of Excellence in Molecular Biology, University of California, San Francisco 94143-0724.

CONTRACT NUMBER: HL-43821 (NHLBI)

SOURCE: MOLECULAR AND CELLULAR BIOLOGY, (1991 Sep) 11 (9) 4627-34.

Journal code: 8109087. ISSN: 0270-7306.

PUB. COUNTRY: United States

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 199109

ENTRY DATE: Entered STN: 19911013

Last Updated on STN: 19970203

Entered Medline: 19910920

AB To determine the mechanisms by which multiple forms of fibroblast growth factor (FGF) receptors are generated, we have mapped the arrangement of exons and introns in the human FGF receptor 1 (FGFR 1) gene (flg). We found three alternative exons encoding a portion of the third immunoglobulin (Ig)-like domain of the receptor. One of these

alternatives

encodes a sequence that is part of a secreted form of FGFR 1. The other two encode sequences that are likely part of transmembrane forms of FGFR 1. One of these forms has not been previously reported in published

cDNAs.

Also, we have determined the structural organization of a portion of the human FGFR 2 gene (bek) and found a similar arrangement of alternative exons for the third Ig-like domain. The arrangement of these genes suggests that there are conserved mechanisms governing the expression of secreted FGF receptors as well as the expression of at least two distinct membrane-spanning forms of the FGF receptors. The diverse forms appear to be generated by alternative splicing of mRNA and selective use of polyadenylation signals.

L14 ANSWER 6 OF 6

MEDLINE

DUPLICATE 3

ACCESSION NUMBER: 97229252 MEDLINE  
DOCUMENT NUMBER: 97229252 PubMed ID: 9075249  
TITLE: Pediatric **brain tumors** express multiple  
receptor tyrosine kinases including novel cell adhesion  
kinases.  
AUTHOR: Weiner H L; Rothman M; Miller D C; Ziff E B  
CORPORATE SOURCE: Department of Neurosurgery (Pediatric Neurosurgery), New  
York University Medical Center, NY 10016, USA.  
CONTRACT NUMBER: P20 NS31088 (NINDS)  
SOURCE: PEDIATRIC NEUROSURGERY, (1996 Aug) 25 (2) 64-71;  
discussion

71-2.

Journal code: 9114967. ISSN: 1016-2291.

PUB. COUNTRY: Switzerland  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 199706  
ENTRY DATE: Entered STN: 19970612  
Last Updated on STN: 20000303  
Entered Medline: 19970603

AB We have used the polymerase chain reaction to clone and characterize  
growth factor receptor tyrosine kinases (RTKs) expressed in 3  
pathologically distinct pediatric **brain tumors**, an  
anaplastic **ependymoma**, a glioblastoma multiforme and a primitive  
neuroectodermal tumor (PNET). These neoplasms are presumed to be derived  
from embryonic neuroepithelial precursor cells of the central nervous  
system. This cloning demonstrated expression of 24 distinct kinase genes:  
16 receptor type kinases and 8 nonreceptor type kinases. The expression  
of 6 receptors, including Hek2, IRR, Ryk, **FGFR3**, and 2 members of  
the newly identified cell adhesion kinase receptor family, DDR and TKT,  
in such tumors has not been reported previously. Northern analysis of mRNA  
levels revealed DDR expression in 6 of 7 pediatric **brain  
tumors** including an **ependymoma**, PNET, glioblastoma and  
**astrocytoma**, and also in an adult pheochromocytoma. Thus, the DDR  
cell adhesion kinase may be widely expressed in pediatric **brain  
tumors**. Also, PCR cloning may be an effective procedure for  
characterizing RTKs in clinical tissue samples and revealing the  
expression of novel RTK species.

WER 5 OF 6 MEDLINE

DUPLICATE 2

ACCESSION NUMBER: 1999085118 MEDLINE  
DOCUMENT NUMBER: 99085118 PubMed ID: 9864407  
TITLE: Fibroblast growth factor-9 (glia-activating factor)  
stimulates proliferation and production of glial

fibrillary

acidic protein in human **gliomas** either in the  
presence or in the absence of the endogenous growth factor  
expression.

AUTHOR: Miyagi N; Kato S; Terasaki M; Aoki T; Sugita Y; Yamaguchi  
M; Shigemori M; Morimatsu M

CORPORATE SOURCE: Department of Pathology, Kurume University, School of  
Medicine, Kurume, Japan.

SOURCE: ONCOLOGY REPORTS, (1999 Jan-Feb) 6 (1) 87-92.  
Journal code: 9422756. ISSN: 1021-335X.

PUB. COUNTRY: Greece

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 199903

ENTRY DATE: Entered STN: 19990402

Last Updated on STN: 19990402

Entered Medline: 19990322

AB We tested fibroblast growth factor-9 (FGF-9) expression in human  
**glioma** cells (U251MG, T98G, U87MG, KALS-1, NMC-G1) and only NMC-G1  
expressed endogenous FGF-9. All cells expressed bFGF and high affinity  
receptors for FGFs (FGFR1 and **FGFR3**). Exogenously supplied bFGF  
and FGF-9 both showed mitogenic activities in all cells. Neutralizing  
antibody against bFGF inhibited the proliferation in U251MG and NMC-G1,  
however that against FGF-9 inhibited the proliferation only in NMC-G1.  
GFAP expression was stimulated by both FGFs in these cells. FGF-9  
potentially regulates proliferation and GFAP expression in human  
**gliomas** either in the presence or in the absence of the endogenous  
growth factor expression.

L14 ANSWER 4 OF 6 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2001:756373 CAPLUS

DOCUMENT NUMBER: 136:51990

TITLE: Expression profiling of **medulloblastoma**:  
PDGFRA and the RAS/MAPK pathway as therapeutic

targets

for metastatic disease

AUTHOR(S): MacDonald, Tobey J.; Brown, Kevin M.; LaFleur,  
Bonnie;

Peterson, Katia; Lawlor, Christopher; Chen, Yidong;  
Packer, Roger J.; Cogen, Philip; Stephan, Dietrich A.  
CORPORATE SOURCE: Center for Cancer and Transplantation Biology,  
Children's National Medical Center, Washington, DC,  
USA

SOURCE: Nature Genetics (2001), 29(2), 143-152

CODEN: NGENEC; ISSN: 1061-4036

PUBLISHER: Nature America Inc.

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Little is known about the genetic regulation of **medulloblastoma** dissemination, but metastatic **medulloblastoma** is highly assocd. with poor outcome. We obtained expression profiles of 23 primary **medulloblastomas** clin. designated as either metastatic (M+) or non-metastatic (M0) and identified 85 genes whose expression differed significantly between classes. Using a class prediction algorithm based on these genes and a leave-one-out approach, we assigned sample class to these tumors (M+ or M0) with 72% accuracy and to four addnl. independent tumors with 100% accuracy. We also assigned the metastatic **medulloblastoma** cell line Daoy to the metastatic class. Notably, platelet-derived growth factor receptor .alpha. (PDGFRA) and members of the downstream RAS/mitogen-activated protein kinase (MAPK) signal transduction pathway are upregulated in M+ tumors. Immunohistochem. validation on an independent set of tumors shows significant overexpression of PDGFRA in M+ tumors compared to M0 tumors. Using in vitro assays, we show that platelet-derived growth factor .alpha. (PDGFA) enhances **medulloblastoma** migration and increases downstream MAP2K1 (MEK1), MAP2K2 (MEK2), MAPK1 (p42 MAPK) and MAPK3 (p44 MAPK) phosphorylation in a dose-dependent manner. Neutralizing antibodies to PDGFRA blocks MAP2K1, MAP2K2 and MAPK1/3 phosphorylation, whereas U0126,

a highly specific inhibitor of MAP2K1 and MAP2K2, also blocks MAPK1/3.

Both inhibit migration and prevent PDGFA-stimulated migration. These results provide the first insight into the genetic regulation of **medulloblastoma** metastasis and are the first to suggest a role for PDGFRA and the RAS/MAPK signaling pathway in **medulloblastoma** metastasis. Inhibitors of PDGFRA and RAS proteins should therefore be considered for investigation as possible novel therapeutic strategies against **medulloblastoma**.

REFERENCE COUNT: 38 THERE ARE 38 CITED REFERENCES AVAILABLE FOR  
THIS

FORMAT RECORD. ALL CITATIONS AVAILABLE IN THE RE

L14 ANSWER 3 OF 6 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
ACCESSION NUMBER: 2001:265659 BIOSIS  
DOCUMENT NUMBER: PREV200100265659  
TITLE: Crystal structure of fibroblast growth factor 9 reveals regions implicated in dimerization and autoinhibition.  
AUTHOR(S): Plotnikov, Alexander N.; Eliseenkova, Anna V.; Ibrahimi, Omar A.; Shriver, Zachary; Sasisekharan, Ram; Lemmon, Mark A.; Mohammadi, Moosa (1)  
CORPORATE SOURCE: (1) Department of Pharmacology, New York University School of Medicine, New York, NY, 10016 USA  
SOURCE: Journal of Biological Chemistry, (February 9, 2001) Vol. 276, No. 6, pp. 4322-4329. print.  
ISSN: 0021-9258.  
DOCUMENT TYPE: Article  
LANGUAGE: English  
SUMMARY LANGUAGE: English

AB Fibroblast growth factors (FGFs) constitute a large family of heparin-binding growth factors with diverse biological activities. FGF9 was originally described as glia-activating factor and is expressed in the nervous system as a potent mitogen for glia cells. Unlike most FGFs, FGF9 forms dimers in solution with a  $K_d$  of 680 nM. To elucidate the molecular mechanism of FGF9 dimerization, the crystal structure of FGF9 was determined at 2.2 Å resolution. FGF9 adopts a beta-trefoil fold similar to other FGFs. However, unlike other FGFs, the N- and C-terminal regions outside the beta-trefoil core in FGF9 are ordered and involved in the formation of a 2-fold crystallographic dimer. A significant surface area (>2000 Å<sup>2</sup>) is buried in the dimer interface that occludes a major receptor binding site of FGF9. Thus, we propose an autoinhibitory mechanism for FGF9 that is dependent on sequences outside of the beta-trefoil core. Moreover, a model is presented providing a molecular basis for the preferential affinity of FGF9 toward **FGFR3**.



L14 ANSWER 2 OF 6 MEDLINE DUPLICATE 1  
 ACCESSION NUMBER: 2001416780 MEDLINE  
 DOCUMENT NUMBER: 21359863 PubMed ID: 11466624  
 TITLE: Frequency of **fibroblast growth factor receptor 3** mutations in sporadic tumours.  
 AUTHOR: Sibley K; Stern P; Knowles M A  
 CORPORATE SOURCE: ICRF Clinical Centre, St. James's University Hospital, Leeds, LS9 7TF, UK.  
 SOURCE: ONCOGENE, (2001 Jul 19) 20 (32) 4416-8.  
 Journal code: 8711562. ISSN: 0950-9232.  
 PUB. COUNTRY: England: United Kingdom  
 DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
 LANGUAGE: English  
 FILE SEGMENT: Priority Journals  
 ENTRY MONTH: 200108  
 ENTRY DATE: Entered STN: 20010813  
 Last Updated on STN: 20010813  
 Entered Medline: 20010809

AB Mutations in **FGFR3** have been identified in several tumour types including bladder carcinoma, cervical carcinoma, and multiple myeloma. In bladder carcinoma, we recently identified **FGFR3** mutations in 41% of tumours, making this the most frequently mutated putative oncogene identified in bladder cancer to date. We have now investigated the frequency of **FGFR3** mutation in a panel of 125 tumours and 13 cell lines from various other organs. We analysed the mutation hotspots

in  
 exons 7, 10 and 15 by direct DNA sequencing, and found one mutation in exon 7 (S249C) in 1/28 (3.5%) cervical tumours. Mutations were not detected in stomach, rectum, colon, prostate, ovarian, breast, **brain**, or renal **tumours**, nor were they found in any of the cell lines included in this study. We conclude that **FGFR3** is commonly mutated in bladder carcinoma and only rarely in cervical carcinoma. Several tumour types appear not to possess any mutations in **FGFR3**, suggesting that these mutations are important only in the development of certain types of tumour.

L20 ANSWER 3 OF 4 MEDLINE DUPLICATE 2  
ACCESSION NUMBER: 1999308632 MEDLINE  
DOCUMENT NUMBER: 99308632 PubMed ID: 10380925  
TITLE: Diverse signaling pathways activated by growth factor  
receptors induce broadly overlapping, rather than  
independent, sets of genes.  
COMMENT: Comment in: Cell. 1999 Jun 11;97(6):675-8  
AUTHOR: Fambrough D; McClure K; Kazlauskas A; **Lander E S**  
CORPORATE SOURCE: Whitehead Institute for Biomedical Research, Cambridge,  
Massachusetts 02142, USA.  
SOURCE: CELL, (1999 Jun 11) 97 (6) 727-41.  
Journal code: 0413066. ISSN: 0092-8674.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 199907  
ENTRY DATE: Entered STN: 19990730  
Last Updated on STN: 20000303  
Entered Medline: 19990722

AB We sought to explore the relationship between receptor tyrosine kinase (RTK) activated signaling pathways and the transcriptional induction of immediate early genes (IEGs). Using global expression monitoring, we identified 66 fibroblast IEGs induced by platelet-derived growth factor beta receptor (PDGFRbeta) signaling. Mutant receptors lacking binding sites for activation of the PLCgamma, PI3K, SHP2, and RasGAP pathways still retain partial ability to induce 64 of these IEGs. Removal of the Grb2-binding site further broadly reduces induction. These results suggest that the diverse pathways exert broadly overlapping effects on IEG induction. Interestingly, a mutant receptor that restores the RasGAP-binding site promotes induction of an independent group of genes, normally induced by interferons. Finally, we compare the PDGFRbeta and **fibroblast growth factor receptor 1**; each induces essentially identical IEGs in fibroblasts.

L14 ANSWER 32 OF 33 PCTFULL COPYRIGHT 2003 Univentio  
 ACCESSION NUMBER: 2001064882 PCTFULL ED 20020822  
 TITLE (ENGLISH): 1983, 52881, 2398, 45449, 50289, AND 52872, G  
 PROTEIN-COUPLED RECEPTORS AND USES THEREFOR  
 TITLE (FRENCH): RECEPTEURS COUPLES A UNE PROTEINE G, NUMEROTEES 1983,  
 52881, 2398, 45449, 50289, ET 52872, ET UTILISATIONS  
 CORRESPONDANTES  
 INVENTOR(S): GLUCKSMANN, Maria, Alexandra; GALVIN, Katherine, M.;  
 SILOS-SANTIAGO, Inmaculada  
 PATENT ASSIGNEE(S): MILLENNIUM PHARMACEUTICALS, INC.; GLUCKSMANN, Maria,  
 Alexandra; GALVIN, Katherine, M.; SILOS-SANTIAGO,  
 Inmaculada  
 DOCUMENT TYPE: Patent  
 PATENT INFORMATION:

	NUMBER	KIND	DATE
	WO 2001064882	A2	20010907
DESIGNATED STATES	AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW GH GM KE LS MW MZ SD SL SZ TZ UG ZW AM AZ BY KG KZ MD RU TJ TM AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR BF BJ CF CG CI CM GA GN GW ML MR NE SN TD TG		
APPLICATION INFO.:	WO 2001-US6543	A	20010228
PRIORITY INFO.:	US 2000-60/186,059		20000229

DETD The subject can be a cancer patient e.g., a patient with **brain cancer**, bone **cancer**, or prostate cancer. In other embodiments, the subject is a non-human animal, e.g., an experimental animal, e.g., an arthritic rat model of. . .

. . . is a method of evaluating a sample. The method includes providing a sample, e.g., from the subject, and determining a gene **expression profile** of the sample, wherein the profile includes a value representing the level of 1983, 52881, 2398, 45449, 50289 and 52872 expression.

The method can further include comparing the value of the gene **expression profile** of the sample to a reference value or reference profile. The gene **expression profile** of the sample can be obtained by any of the methods described herein.

15 (e.g., by providing a nucleic acid. . . an indication that the subject has or is disposed to having a disorder as described herein. The method can be used to **monitor** a **treatment** for such disorders in a subject. For example, the gene **expression profile** can be determined for a sample from a subject undergoing **treatment**.

L14 ANSWER 22 OF 33 PCTFULL COPYRIGHT 2003 Univentio  
 ACCESSION NUMBER: 2001083781 PCTFULL ED 20020826  
 TITLE (ENGLISH): 14094, A NOVEL HUMAN TRYPSIN FAMILY MEMBER AND USES  
 THEREOF  
 TITLE (FRENCH): 14094, UN NOUVEAU MEMBRE DANS LA FAMILLE DE LA  
 TRYPSINE HUMAINE ET SON UTILISATION  
 INVENTOR(S): MEYERS, Rachel; MACBETH, Kyle, J.  
 PATENT ASSIGNEE(S): MILLENNIUM PHARMACEUTICALS, INC.; MEYERS, Rachel;  
 MACBETH, Kyle, J.  
 DOCUMENT TYPE: Patent  
 PATENT INFORMATION:

	NUMBER	KIND	DATE
	WO 2001083781	A2	20011108
DESIGNATED STATES	AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW GH GM KE LS MW MZ SD SL SZ TZ UG ZW AM AZ BY KG KZ MD RU TJ TM AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR BF BJ CF CG CI CM GA GN GW ML MR NE SN TD TG		
APPLICATION INFO.:	WO 2001-US13903	A	20010430
PRIORITY INFO.:	US 2000-60/200,621		20000428
	US 2000-09/633,300		20000808

DETD . . . cancerous or pre-cancerous tissue where a 14094 polypeptide or nucleic acid is expressed, e.g., breast, ovarian, colon, liver, lung, kidney, or **brain cancer**.

. . . found in a tissue 1, where a 14094 polypeptide or nucleic acid is expressed, e.g., breast, ovarian, colon, liver, lung, kidney, or **brain cancer**.

. . . cancer is a sarcoma, a carcinoma, or an adenocarcinoma. Preferably, the cancer is a breast, ovarian, colon, lung, liver, kidney, or **brain cancer**.

. . . gastric cancer, esophageal cancer, rectal cancer, pancreatic cancer, ovarian cancer, prostate cancer, uterine cancer, cancer of the head and neck, skin **cancer**, **brain cancer**, squamous cell carcinoma, sebaceous gland carcinoma.

. . . can further include comparing the value or the profile (i.e., multiple values) to a reference value or reference profile. The gene **expression profile** of the sample can be obtained by any of the methods described herein (e.g., by providing a nucleic acid from the sample. . . is an indication that the subject has or is disposed to having a cell proliferative disorder. The method can be used to **monitor** a **treatment** for a cell proliferative disorder in a subject. For example, the gene **expression profile** can be

determined for a sample from a subject

75 -

undergoing **treatment**. The profile can be compared to a reference profile or to a profile obtained from the subject prior to **treatment** or prior to onset of the disorder (see, e.g., Golub et al. (1999) Science 286:53 I).

context,  
the effect of one cell type on another cell type in response to a biological stimulus can be determined, e.g., to **monitor** the effect of cell-cell interaction at the level of gene expression, In another embodiment, cells are contacted with a **therapeutic** agent. The **expression profile** of the cells is determined using the array, and the **expression profile** is compared to the profile of like cells not contacted with the agent.

For

example, the assay can be used to determine or analyze the molecular basis of an undesirable effect of the **therapeutic** agent. If an agent is administered **therapeutically** to treat one cell type but has an undesirable effect on another cell type, the invention provides an

assay

to determine. . .

L8 ANSWER 12 OF 20 MEDLINE

ACCESSION NUMBER: 2001393249 MEDLINE

DOCUMENT NUMBER: 21064203 PubMed ID: 11122874

TITLE: Application of advances in molecular biology to the  
**treatment** of brain tumors.

AUTHOR: Takeshima H; Sawamura Y; Gilbert M R; Van Meir E G

CORPORATE SOURCE: Department of Neurosurgery, Faculty of Medicine, Kagoshima  
University, 8-35-1 Sakuraga-oka, Kagoshima 890-8520,  
Japan.. m2040k@khosp2.kufm.kagoshima-u.ac.jp

SOURCE: Curr Oncol Rep, (2000 Sep) 2 (5) 425-33. Ref: 56  
Journal code: 100888967. ISSN: 1523-3790.

PUB. COUNTRY: United States

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
General Review; (REVIEW)  
(REVIEW, TUTORIAL)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 200107

ENTRY DATE: Entered STN: 20010716

Last Updated on STN: 20010716

Entered Medline: 20010712

AB Recent advances in molecular biology have substantially improved our understanding of the molecular genetics of primary brain neoplasms. Soon each histopathologic category of glioma will be further divided into subgroups according to similar genetic background, gene **expression profile**, and similarity of biologic responses to radiotherapy or chemotherapy. Identification of key molecules that are specifically altered in neoplastic cells will provide candidate molecular targets for tumor **treatment**. Novel **therapeutic** tools for targeting tumor cells, such as viral vectors for gene **therapy**, have been created. In the near future, the accumulation of new knowledge in brain tumor biology and genetics, combined with rational drug design, will revolutionize the **treatment** of malignant gliomas, which are among the most lethal human cancers.

L14 ANSWER 3 OF 33 PCTFULL COPYRIGHT 2003 Univentio  
 ACCESSION NUMBER: 2002059610 PCTFULL ED 20020809 EW 200231  
 TITLE (ENGLISH): USING OVEREXPRESSION OF LAMININ ALPHA 4 SUBUNIT AS A  
 DIAGNOSTIC AND PROGNOSTIC INDICATOR OF MALIGNANT  
 TUMORS  
 TITLE (FRENCH): UTILISATION DE LA SUREXPRESSION DE LA SOUS-UNITE DE  
 LAMININE ALPHA 4 EN TANT QU'INDICATEUR DIAGNOSTIQUE ET  
 PRONOSTIQUE DE TUMEURS MALIGNES  
 INVENTOR(S): LJUBIMOVA, Julia, Y.; LJUBIMOV, Alexander, V.; BLACK,  
 Keith, L.  
 PATENT ASSIGNEE(S): CEDARS-SINAI MEDICAL CENTER  
 AGENT: STEINBERG, Nisan, A., Ph.D.  
 LANGUAGE OF FILING: English  
 LANGUAGE OF PUBL.: English  
 DOCUMENT TYPE: Patent  
 PATENT INFORMATION:

NUMBER	KIND	DATE
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DESIGNATED STATES	WO 2002059610	A2 20020801
	AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR	
	CU CZ DE DK DM DZ EC EE ES FI GB GD GE GH GM HR HU ID	
	IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD	
	MG MK MN MW MX MZ NO NZ PH PL PT RO RU SD SE SG SI SK	
	SL TJ TM TR TT TZ UA UG UZ VN YU ZA ZW GH GM KE LS MW	
	MZ SD SL SZ TZ UG ZM ZW AM AZ BY KG KZ MD RU TJ TM AT	
	BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR	
	BF BJ CF CG CI CM GA GN GW ML MR NE SN TD TG	

APPLICATION INFO.: WO 2001-US50292 A 20011219  
 PRIORITY INFO.: US 2000-09/741,550 20001219

ABEN Disclosed is a method of diagnosing the presence of a malignant tumor,  
 including a glioma, in a human subject, which involves detecting  
 overexpression of laminin \*4 subunit protein or laminin \*4-specific  
 mRNA, compared to the expression level in a normal tissue control. Also  
 disclosed are a method of predicting the recurrence of a malignant  
 tumor  
 in a human subject from whom a malignant tumor has been resected and a  
 method of classifying the grade of a malignant tumor, such as a glial  
 tumor, based on a molecular classification.  
 ABFR L'invention concerne un procede de diagnostic de la presence de tumeurs  
 malignes, y compris de gliomes, chez l'homme, qui implique la detection  
 de la surexpression de la proteine de sous-unite de laminine &alpha;4  
 ou  
 d'un ARNm specifique de la laminine &alpha;4, par comparaison avec le  
 niveau d'expression dans un temoin tissulaire normal. L'invention  
 concerne egalement un procede permettant de prevoir la recurrence d'une  
 tumeur maligne chez un homme ayant subi une resection de tumeur  
 maligne,  
 ainsi qu'un procede de determination de la classe d'une tumeur maligne,  
 telle qu'une tumeur gliale, sur la base d'une classification  
 moleculaire.

L14 ANSWER 4 OF 33 PCTFULL COPYRIGHT 2003 Univentio  
 ACCESSION NUMBER: 2002057457 PCTFULL ED 20020801 EW 200230  
 TITLE (ENGLISH): 55562 AND 21617, NOVEL HUMAN PROTEINS AND METHODS OF  
 USE THEREOF  
 TITLE (FRENCH): 55562 ET 21617, NOUVELLES PROTEINES HUMAINES ET LEURS  
 METHODES D'UTILISATION  
 INVENTOR(S): MEYERS, Rachel, A.; BANDARU, Rajasekhar  
 PATENT ASSIGNEE(S): MILLENNIUM PHARMACEUTICALS, INC., for all designates  
 States except US; MEYERS, Rachel, A., for US only;  
 BANDARU, Rajasekhar, for US only  
 AGENT: COLLAZO, Diana, M.  
 LANGUAGE OF FILING: English  
 LANGUAGE OF PUBL.: English  
 DOCUMENT TYPE: Patent  
 PATENT INFORMATION:

	NUMBER	KIND	DATE
	WO 2002057457	A2	20020725
DESIGNATED STATES	AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK DM DZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ OM PH PL PT RO RU SD SE SG SI SK SL TJ TM TN TR TT TZ UA UG US UZ VN YU ZA ZM ZW GH GM KE LS MW MZ SD SL SZ TZ UG ZM ZW AM AZ BY KG KZ MD RU TJ TM AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR BF BJ CF CG CI CM GA GN GQ GW ML MR NE SN TD TG		

APPLICATION INFO.: WO 2001-US49416 A 20011218  
 PRIORITY INFO.: US 2000-60/256,249 20001218  
 US 2000-60/256,405 20001218

ABEN The invention provides isolated nucleic acids molecules, designated  
 21617 and 55562 nucleic acid molecules, which encode novel  
 dehydrogenase  
 or tetratricopeptide repeat members. The invention also provides  
 antisense nucleic acid molecules, recombinant expression vectors  
 containing 21617 or 55562 nucleic acid molecules, host cells into which  
 the expression vectors have been introduced, and nonhuman transgenic  
 animals in which a 21617 or 55562 gene has been introduced or  
 disrupted.

The invention still further provides isolated 21617 or 55562 proteins,  
 fusion proteins, antigenic peptides and anti-21617 or 55562 antibodies.  
 Diagnostics methods utilizing compositions of the invention are also  
 provided.

ABFR La presente invention concerne des molecules d'acides nucleiques  
 isolees, designees sous le nom de molecules d'acides nucleiques 21617  
 et

55562, codant de nouvelles repetitions de la deshydrogenase ou du  
 tetratricopeptide. L'invention concerne egalement des molecules  
 d'acides nucleiques antisens, des vecteurs d'expression recombinés  
 contenant ces molecules d'acides nucleiques 21617 ou 55562, des  
 cellules  
 hotes dans lesquelles ces vecteurs d'expression ont ete introduits et  
 des animaux transgeniques non humains dans lesquels le gene 21617 ou  
 55562 a ete introduit ou interrompu. L'invention concerne egalement des  
 proteines 21617 ou 55562 isolees, des proteines hybrides, des peptides  
 antigeniques et des anticorps anti-21617 ou 55562. L'invention concerne  
 egalement des methodes diagnostiques utilisant des compositions de la



presente invention.

L2 ANSWER 11 OF 23 SCISEARCH COPYRIGHT 2003 ISI (R)  
ACCESSION NUMBER: 2000:49185 SCISEARCH  
THE GENUINE ARTICLE: 273BX  
TITLE: Cell-type specific expression in the pituitary:  
physiology  
and gene therapy  
AUTHOR: Castro M G (Reprint); Windeatt S; SmithArica J;  
Lowenstein  
CORPORATE SOURCE: P R  
UNIV MANCHESTER, SCH MED, MOL MED UNIT, ROOM I-302,  
OXFORD  
RD, MANCHESTER M13 9PT, LANCs, ENGLAND (Reprint)  
COUNTRY OF AUTHOR: ENGLAND  
SOURCE: BIOCHEMICAL SOCIETY TRANSACTIONS, (DEC 1999)  
Vol. 27, Part 6, pp. 858-863.  
Publisher: PORTLAND PRESS, 59 PORTLAND PLACE, LONDON W1N  
3AJ, ENGLAND.  
ISSN: 0300-5127.  
DOCUMENT TYPE: Article; Journal  
FILE SEGMENT: LIFE  
LANGUAGE: English  
REFERENCE COUNT: 49

L2 ANSWER 18 OF 23 SCISEARCH COPYRIGHT 2003 ISI (R)  
ACCESSION NUMBER: 1998:946863 SCISEARCH  
THE GENUINE ARTICLE: 146TR  
TITLE: The cytogenesis and pathogenesis of pituitary adenomas  
AUTHOR: Asa S L (Reprint); Ezzat S  
CORPORATE SOURCE: MT SINAI HOSP, DEPT PATHOL & LAB MED, 600 UNIV AVE,  
TORONTO, ON M5G 1X5, CANADA (Reprint); MT SINAI HOSP,  
DEPT  
MED, TORONTO, ON M5G 1X5, CANADA; UNIV TORONTO, DEPT LAB  
MED & PATHOBIOL, TORONTO, ON M5G 1X5, CANADA; UNIV  
TORONTO, DEPT MED, TORONTO, ON M5G 1X5, CANADA  
COUNTRY OF AUTHOR: CANADA  
SOURCE: ENDOCRINE REVIEWS, (DEC 1998) Vol. 19, No. 6,  
pp. 798-827.  
Publisher: ENDOCRINE SOC, 4350 EAST WEST HIGHWAY SUITE  
500, BETHESDA, MD 20814-4110.  
ISSN: 0163-769X.  
DOCUMENT TYPE: General Review; Journal  
FILE SEGMENT: LIFE  
LANGUAGE: English  
REFERENCE COUNT: 361

L2 ANSWER 16 OF 23 SCISEARCH COPYRIGHT 2003 ISI (R)

ACCESSION NUMBER: 1999:271844 SCISEARCH

THE GENUINE ARTICLE: 182EY

TITLE: The pathology of pituitary tumors

AUTHOR: Asa S L (Reprint)

CORPORATE SOURCE: MT SINAI HOSP, DEPT PATHOL & LAB MED, 600 UNIV AVE,  
TORONTO, ON M5G 1X5, CANADA (Reprint); UNIV TORONTO, DEPT  
LAB MED & PATHOBIOL, TORONTO, ON, CANADA

COUNTRY OF AUTHOR: CANADA

SOURCE: ENDOCRINOLOGY AND METABOLISM CLINICS OF NORTH AMERICA, (  
**MAR 1999**) Vol. 28, No. 1, pp. 13-&.

Publisher: W B SAUNDERS CO, INDEPENDENCE SQUARE WEST  
CURTIS CENTER, STE 300, PHILADELPHIA, PA 19106-3399.

ISSN: 0889-8529.

DOCUMENT TYPE: General Review; Journal

FILE SEGMENT: LIFE; CLIN

LANGUAGE: English

REFERENCE COUNT: 153

\*ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS\*

AB The pathologist plays an important role in the distinction of  
pituitary

adenomas from other tumors and tumor-like lesions of the sellar region,  
and in the accurate morphologic characterization of pituitary adenomas. A  
clinicopathologic classification of pituitary adenomas is based on cell  
differentiation correlated with clinical evidence of hormone secretion;  
this classification emphasizes clinically relevant features that can

offer

guidance for patient management. The application of a rational approach

to

the immunohistochemical analysis of these lesions can be used to evaluate  
pathogenetic and prognostic markers and to predict responses to specific  
therapeutic modalities.